

Tolerances & Evaluation Criteria (2 "tier approach")

Wanted to state minimum acceptable tolerance for TPS "basic" dose calculation:

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- "The tolerances for the basic photon tests are widely accepted as minimum criteria for static photon beams under conditions of charged particle equilibrium."
- Wanted to push the limit on some evaluation criteria to find limitations of dose calculations:
 - "Given that there is not widely accepted minimum tolerance for the other verification tests in this MPPG, (including those for VMAT/IMRT), those evaluation criteria must not be interpreted as mandatory or regulatory tolerances. Rather, they are values defined as points for further investigation, possible improvement, and resolution."
- Did not want to state or use any minimum tolerance values not widely accepted/published: - "All the tolerances and criteria in this report are based on a combinat
 - "All the tolerances and criteria in this report are based on a combination of published guidelines, the dosimetric audits performed by the Radiological Physics Center, and the experience of authors. Users are encouraged to not only meet these tolerances, but also strive to achieve dosimetric agreement comparable to that reported in the literature for their particular algorithm."

Goals

While the implementation of robust and comprehensive QA programs recommended in other AAPM reports is strongly encouraged, the overall objective of this MPPG is to provide an overview of the minimum requirements for TPS dose algorithm commissioning and QA in a clinical setting. Specific goals for this report are to:

- Clearly identify and reference applicable portions of existing AAPM reports and peer-reviewed articles for established commissioning components.
- Provide updated guidelines on technologies that have emerged since the
- Provide guidance on validation tests for dose accuracy and constancy (select downloadable datasets/contours & beam parameters are provided for optional
- downloadable datasets/contours & beam parameters are provided for optiona use).
- Provide guidance on <u>typical achievable tolerances and evaluation criteria for</u> <u>clinical implementation</u>.
- Provide a checklist for commissioning processes and associated documentation.

Scope/exclusions

- Title: Commissioning and QA of Treatment Planning <u>Dose</u> Calculations: Megavoltage Photon and Electron Beams
- The scope of this report is limited to the commissioning and QA of the beam modeling and calculation portion of a TPS where:
 - External photon and electron treatment beams are delivered at typical SSDs using a gantry mounted radiation source including conventional and small fields used in IMRT, VMAT, helical tomotherapy delivery, and SRS/SBRT (still up for discussion).
- Modern dose algorithms are utilized including corrections for tissue heterogeneity.
 The Multi-Leaf Collimator (MLC) is used as the primary method of shaping the beam aperture for treatments. (individually fabricated IMRT modifiers, cones... still up for discussion)
- Excludes: (not an exhaustive list, and not all written in document)
 Non-dosimetric components of system, e.g.: DVH, leaf sequences, contours, image
 registration...
 - Brachytherapy
 - Proton therapy
 - Non-commercial planning systems
 - Radiation delivered by robots

(plus rati	Outline: follow outline of MPPG onale & some implementation experiences)
1.	Introduction
	a. Goals
	 Tolerances and evaluation criteria
	c. Scope/exclusions
2.	Staff qualifications
3.	Data acquisition
4.	Model within TPS software
5.	Photon beams: basic dose algorithm validation
	 MatLab code for 1D gamma analysis
	 Trilogy: absolute dose verification, large field/off axis MLC tests
	 TomoTherapy: "tomophants"
6.	Photon beams: heterogeneity correction validation
	 Clinac: CIRMS phantom
7.	Photon beams: IMRT/VMAT dose validation
	 TomoTherapy – TG 119 tests and clinical case
8.	Electron beams
9.	Routine QA (downloadable datasets)

Data Acquisition Question

What data do you use when commissioning a new dose algorithm?

- 1. Collect data according to vendors guidelines
- 2. Collect some of the vendor recommended data but not all
- 3. Collect all required data and more
- 4. Use golden beam data
- 5. Hey, I thought this wasn't a SAM session.

Staff, Data, Model...

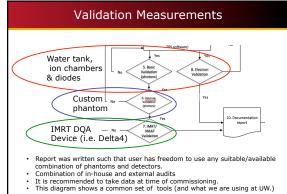
- Staff qualifications QMP, defer to supervision MPPG
- Data acquisition defer to TPS manuals for all required data (water
- tank, and in air for MC) & refer to TG 106. An equipment list/ summary on small field/MLC data acquisition is included:
- PDD and OF with a small volume detector down to at least 2x2 cm² MLC intra and inter-leaf transmission and leaf gap:
- · Large chamber if an average intra- and inter-leaf value is
 - specified. • Separate measurements, use small chamber under the leaf
 - and film for inter-leaf leakage measurements
- Measure leaf-end penumbra with a small detector (such as a diode or micro-chamber) to avoid volume-averaging effects - Leaf timing for binary MLC systems should be verified using film or
- exit detector measurements Model - refer to manual, iterate as needed using results from
- validation testing

Outline: follow outline of MPPG (plus rationale & some implementation experiences) Introduction Goals b. Tolerances and evaluation criteria Scope/exclusions Staff qualifications Data acquisition Model within TPS software Photon beams: basic dose algorithm validation MatLab code for 1D gamma analysis Trilogy: absolute dose verification, large field/off axis MLC tests TomoTherapy: "tomophants" Photon beams: heterogeneity correction validation 6. Clinac: CIRMS phantom Photon beams: IMRT/VMAT dose validation TomoTherapy - TG 119 tests and clinical case Electron beam validation

Validation Question

What type of dose algorithm validation do you do as part of the commissioning process?

- 1. None
- 2. Routine patient specific DQA serves as validation
- 3. In-house test suite (chamber, array, films etc...)
- 4. Peer review audit (colleague or RPC)
- 5. Combination of 3 and 4



5. Basic Validation: Photon beams

Section 5 (Photons in homogeneous media) has 2 sets

- of tests:
 5.1-5.3: "sanity check" of commission data → physics module → planning module and TG 51 calibration value
 - 5.4-5.9: test fields that were not used in commissioning. Compare measured and calculated dose distribution.
 - Tests should be run for each unique configured beam (energy and wedge)

est	Comparison	Description	Tolerance
.1	Dose distributions in planning module vs. modeling (physics) module	Comparison of dose distribution for large (>30x30) field.	Identical
	Dose in test plan vs. clinical calibration condition*	Reference calibration condition check	0.5%
.3	Dose distribution calculated in planning system vs. commissioning data	PDD and off axis factors for a large and a small field size	2%
	dditional measurements requir	red for these tests the necessarily of linac per TG 51	

Varian 2100s		inini materi be	etween tw	
10 MV beams	Meas.(Gy)	TPS calc (Gy)	% diff	1 /\ I
Open, 90 cm SSD	0.893	0.891	-0.18	90 cm SSD
15° W, 90 cm SSD	0.669	0.669	-0.01	
30° W, 90 cm SSD	0.543	0.544	0.21	1 / /
45° W, 90 cm SSD	0.470	0.473	0.71	
60° W, 90 cm SSD	0.392	0.394	0.42	
Open 100 cm SSD	0,744	0.741	-0.34	
The 10 MV 45° wedge exe and is being investigated	ceeded the 0.5% 1	tolerance suggested i	n the MPPG	

Test	Description	Sample tests from literature [7]
5.4	Small MLC shaped field (non SRS)	Photon Test 1
5.5	Large MLC shaped field with extensive blocking (e.g.: mantle)	Photon Test 3
5.6	Off-axis MLC shaped field, with maximum allowed leaf over travel.	Photon Test 2
5.7	Asymmetric MLC shaped field at minimal anticipated SSD	Photon Test 6
5.8	MLC shaped field at oblique incidence (30°)	Photon Test 10
5.9	Large (>15cm) MLC field for each a non-physical wedge angle**	
sts 5 cal v	: high dose, penumbra, and low dose tail reg .4-5.8 are intended for each open and (hard) vedges are considered an extension of the co spectra and only require the addition of Test !) wedged field. Non- rresponding open fie

Accuracy question

How accurate is your worst off axis relative dose calculation?

- 1.1%
- 2.2%
- 3.3%
- 4.4%
- 5.5%

Accuracy question 2

How accurate is your worst off axis relative dose calculation with a wedge in place?

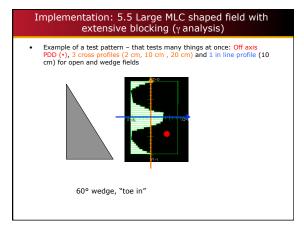
- 1.1%
- 2.2%
- 3.3%
- 4.4%
- 5.5%

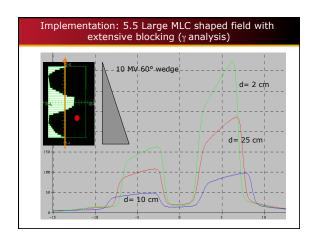
Section 5	: Basic	photon	tolerances
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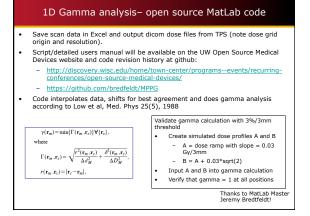
Table 5. Basic TPS photon beam evaluation methods and tolerances

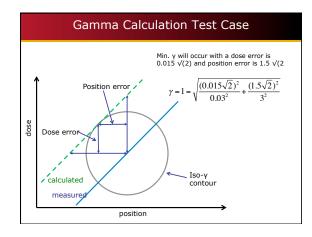
Region	Evaluation Method	Tolerance* (consistent with RPC)
High dose	Relative dose with one parameter change from reference conditions	2%
	Relative dose with multiple parameter changes **	5%
Penumbra	Distance to agreement	3 mm
Low dose tail	Up to 5 cm from field edge	3% of maximum field dose

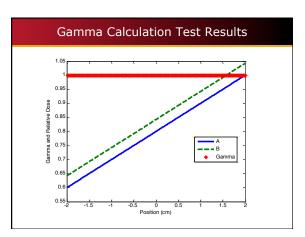
* Tolerances are relative to local dose unless otherwise noted. **e.g.: off axis with physical wedge.

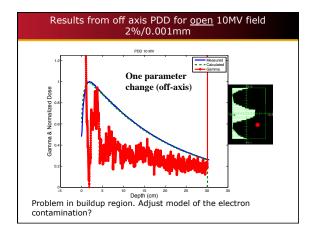


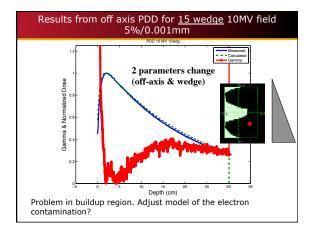


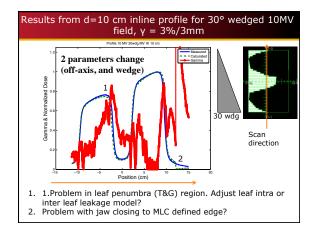


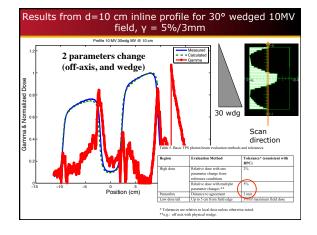


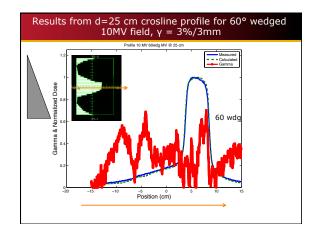


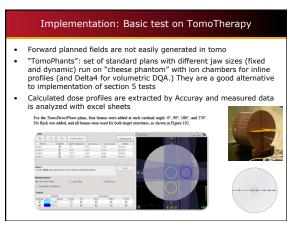


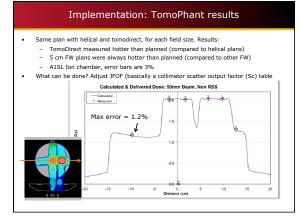










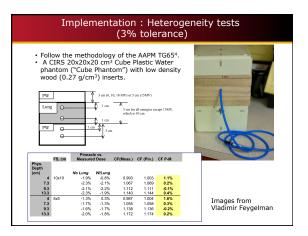


Heterogeneity questions

Which algorithm is not acceptable for dose calculation for lung?

- 1. Pencil beam
- 2. Monte Carlo
- 3. Convolution superposition
- 4. Discreet ordinance (grid based Boltzman solver)
- 5. All are acceptable

	6: Heterogeneous TPS photo		Tolerances*	Reference
Test	Objective	Description	1 oterances*	
6.1	Validate planning system	CT-density calibration for air,		TG 65 [23]; IAEA
	reported electron (or mass)	lung, water, dense bone, and		TRS-430 [7]
	densities against known	possibly additional tissue types.		
	values.			
6.2	Heterogeneity correction	5x5 cm2, measure dose ratio	3%	Carrasco et al. [52
	distal and proximal to lung	above and below heterogeneity		
	tissue	outside of the buildup region		
* Tole		se unless otherwise noted. thms (C/S. MC, GBBS, 1 and heterogeneity (not		ındaries,



`est	Objective	Description (example)	Detector
7.1	Verify small field PDD	$\ge 2x2 \text{ cm}^2 \text{ MLC}$ shaped field, with	Diode or plastic
		PDD acquired at a clinically relevant SSD.	scintillator
7.2	Verify output for small MLC-	Use small square and rectangular	Diode, plastic
	defined fields	MLC-defined segments, measuring	scintillator, mini-
		output at a clinically relevant depth for	chamber or micro-
		each*	ion chamber
7.3	TG-119 tests	Plan, measure, and compare planning	
		and QA results to the TG119 report for	
		both the Head and Neck and C-shape	
		cases.	
7.4	Clinical tests	Choose at least 2 relevant clinical	Ion chamber, film
		cases. Plan, measure, and perform an	and/or array
		in-depth analysis of the results.	
7.5	External review	Simulate, plan, and treat an	Various options
		anthropomorphic phantom with	exist.**
		embedded dosimeters.	

Section 7: IMRT/VMAT Verification

IMRT DQA Question 1

What gamma criteria do you use for patient specific delivery QA (DQA)?

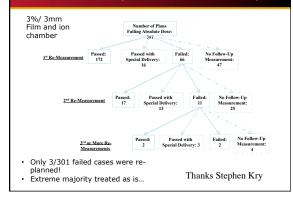
- 1. 1%/1mm
- 2. 2%/2mm
- 3. 3%/3mm
- 4.4%/4mm
- 5. I don't do patient specific DQA and/or I don't use gamma criteria for DQA analysis.

IMRT DQA Question 2

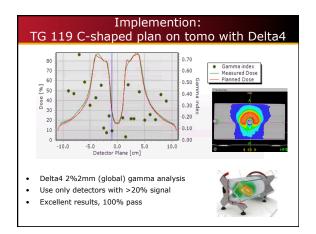
What do you do when a case 'fails' that criteria?

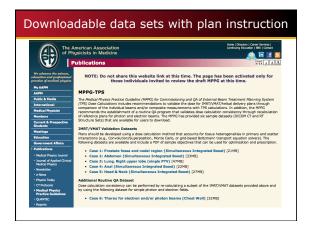
- 1. increase tolerance by 1%/1mm
- 2. Re-measure
- 3. Re-plan
- 4. Pick tolerance so >95% pass and report tolerance values
- 5. My plans never fail

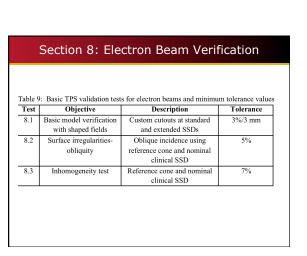
MD Anderson Experience with failed DQA's



able	7: VMAT/IMRT Test Sun	nmary.					
Fest	Objective	De	scription (example)	D	etector	Ref	
7.1	Verify small field PDD		cm ² MLC shaped field,		or plastic	TG-155 (to be	
			h PDD acquired at a	sci	ntillator	published in	
			ically relevant SSD.			MP)	
7.2	Verify output for small	Use small square and rectangular		Diode, plastic		Cadman et al.	
	MLC-defined fields				ntillator,	[53]	
			ing output at a clinically		hamber or		C-shape plan, or
		rele	vant depth for each*		cro-ion		
7.3	TG-119 tests	DI	measure, and compare	ct	amber	TG-1191311	tomo
1.5	1G-119 tests		ng and OA results to the			16-119[31]	Transverse
			report for both the Head				T
			eck and C-shane cases.			IJ	
7.4	Clinical tests		ose at least 2 relevant	Ion	chamber.	Nelms et al.	
		clinica	l cases. Plan, measure,	film a	nd/or array	[54]	
		and pert	form an in-depth analysis				
			of the results.				
7.5	External review		Simulate, plan, and treat an		us options	Kry et al. [32]	
		anthrop	omorphic phantom with	c:	cist.**		₩ 4 65 ►
		en	Table 8: VMAT/IMRT E			4.7.1	
			Measurement Method	valuation	Region	nd Toterances	Tolerance
			Ion Chamber			ent target region	2% of prescribed dose
			ton comment		OAR regi		3% of prescribed dose
Planar/Volumetric Arra				All regions			2%/2mm*, no pass rate
							tolerance, but areas that do not
							pass need to be investigated
End-to-End					I and and	ent target region	5% of prescribed dose







Section 9 QA

- Annually or after major TPS upgrades
- Reference plans should be selected at the time of commissioning and then re-calculated for routine QA comparison. •
- calculated for routine QA comparison. For photons, representative plans for each configured beam should be chosen from Table 4 for static and wedge beams and Table 7 for IMRT/VMAT. Optionally, an additional thorax dataset with contours and suggested static beam parameters can be downloaded and used for some of these tests, (http://www.aapm.org/pubs/tg244/). A 10x10 cm² field and a small field (e.g. 5x5 cm²) can be prescribed to the isocenter located in the center of the PTV. Wedge fields and dynamic arc plans can also be calculated on the thorax data set.
- For electrons, plans should be calculated for each energy using a heterogeneous dataset with reasonable surface curvature. The sample thorax dataset is also suitable for this test. Recommended plans also include extended distance and bolus verification. •
- The routine QA re-calculation should agree with the reference dose calculation to within 1%/1mm. A complete re-commissioning (including validation) may be required if more significant deviations are observed.

	TG244	TG244 Item	Commission
	Section		Report Page
	1	QMP understands algorithms and has received proper	
		training.	_
	3	Manufacturer's guidance for data acquisition was consulted	
Checklist to		and followed.	
quide	3.b	Appropriate CT calibration data acquired.	
guiue	3.d	Review of raw data (compare with published data, check for	
commissioning		error, confirm import into TPS).	
	4	Beam modeling process completed according to vendor's	
report		instructions.	
	4	Beam models evaluated qualitatively and quantitatively using	
		metrics within the modeling software.	
	5	For each beam model perform validation tests 5.1-5.8 (5.9	
		tolerances in Tables 3 and 4.	
	6	Heterogeneity corrections validated for photon beams	
		according to Table 6.	
	7	IMRT and VMAT validations accomplished for each	
		configured beam according to tests 7.1-7.4 in Table 7.	
	7	End-to-End test with external review accomplished for IMRT	
		and VMAT (test 7.5 in Table 7).	
	7	Understand and document limitations of IMRT/VMAT	
		modeling and dose algorithms.	
	8	Electron validations performed according to tests 8.1-8.3 in	
		Table 9.	
	9	Baseline QA plan(s) (for model constancy) identified for each	
		configured beam and routine QA established.	
	10	Peer review obtained and any recommendations addressed.	

Next steps....

- Respond to public comment reviewer comments
- Submit to JACMP - await final review
- Continue implementation of MPPG on Varian, TomoTherapy and • Elekta (AAPM annual meeting abstract)
 - Fine tune gamma analysis in MatLab code, analyze remaining Trilogy and Infinity data - Take heterogeneous and electron data
 - _
 - Create test suite for each machine type (Pinnacle/Eclipse plans, R&V entry and scan Q's)
- . Make gamma analysis code easily available (and easier data input)

Thanks to my collaborators, and to you for your attention!

- All MPPG#5 members!
- UW clinical physicists who helped with implementation
 - Adam Bayliss
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